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STRUCTURE FILE UPDATES: 25 AUG 2011 HIGHEST RN 1323485-64-8 DICTIONARY FILE UPDATES: 25 AUG 2011 HIGHEST RN 1323485-64-8

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http://www.cas.org/support/stngen/stndoc/properties.html

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exact/norm bonds :
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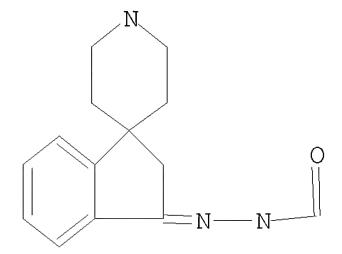
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L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

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SEARCH TIME: 00.00.01

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100.0% PROCESSED 430 ITERATIONS 41 ANSWERS

SEARCH TIME: 00.00.01

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This file contains CAS Registry Numbers for easy and accurate substance identification.

26/08/2011

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L4 1 L3

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L4 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2011 ACS on STN GI

$$R^{8}$$
 R^{9}
 R^{9

Title compds. I [X = O, amino; Y = bond, CO, CS, SOO-2; R1 = H, alkyl, alkoxycarbonyl, etc.; R2-3 = H, halo, CN, alkyl, etc.; R4 = halo, NO2, CN, etc.; Ra = H, halo, OH, CN, etc.; p, q = 0-6; R8 = alk(en/yn)yl, etc.] are prepared For instance, II is prepared in 3 steps from spiro[indan-1-one-3,4'-piperidine]-1'-carboxylic acid tert Bu ester, 4-chlorocinnamyl chloride and hydroxylamine (E (dominant) and Z oximes isolated). Selected example compds. gave >80% control of Spodoptera littoralis. I are useful in controlling insects, acarines, nematodes or molluscs.

AN 2005:570877 CAPLUS

DN 143:77964

TI Preparation of insecticidal spiroindane derivatives

IN Cassayre, Jerome; Molleyres, Louis-Pierre; Maienfisch, Peter; Cederbaum, Fredrik

PA Syngenta Participations A.-G., Switz.

SO PCT Int. Appl., 114 pp. CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

	PATENT NO.						D	DATE			APPL	DATE								
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ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT
     CASREACT 143:77964; MARPAT 143:77964
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     RL: AGR (Agricultural use); BSU (Biological study, unclassified); RCT
     (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (preparation of insecticidal spiroindane derivs. as insecticides,
        acaracides, molluscicides and nematocides)
     855849-47-7 CAPLUS
RN
     Spiro[1H-indene-1, 4'-piperidine]-1'-carboxylic acid,
CN
     3-[2-[(2-chloro-4-pyridinyl)carbonyl]hydrazinylidene]-2,3-dihydro-,
     1,1-dimethylethyl ester (CA INDEX NAME)
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RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of insecticidal spiroindane derivs. as insecticides, acaracides, molluscicides and nematocides)

RN 855849-48-8 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-,

2-[2,3-dihydro-1'-[[4-(trifluoromethoxy)phenyl]methyl]spiro[1H-indene-1,4'-

piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-49-9 CAPLUS
CN 4-Pyridinecarboxylic acid, 2-chloro-,
2-[2,3-dihydro-1'-[[4-[[(1-mothydlerine(14-indexe-1,4'-niperidin

methylethoxy)carbonyl]amino]phenyl]methyl]spiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-50-2 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[[4-(2-ethyl-2H-tetrazol-5-yl)phenyl]methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-51-3 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(4-cyanophenyl)methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-52-4 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(4-fluorophenyl)methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-53-5 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[2,3-dihydro-1'-(phenylmethyl)spiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-54-6 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(2,6-difluorophenyl)methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-55-7 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(3-chlorophenyl)methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-56-8 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[2,3-dihydro-1'-(1-phenylethyl)spiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-57-9 CAPLUS
CN 4-Pyridinecarboxylic acid, 2-chloro-,
2-(2,3-dihydro-1'-methylspiro[1H-indene-1,4'-piperidin]-3ylidene)hydrazide (CA INDEX NAME)

RN 855849-58-0 CAPLUS
CN Spiro[1H-indene-1,4'-piperidine]-1'-carboxylic acid,
6-chloro-3-[2-[(2-chloro-4-pyridinyl)carbonyl]hydrazinylidene]-2,3-dihydro, 1,1-dimethylethyl ester (CA INDEX NAME)

RN 855849-59-1 CAPLUS
CN Spiro[1H-indene-1,4'-piperidine]-1'-carboxylic acid,
5-chloro-3-[2-[(2-chloro-4-pyridinyl)carbonyl]hydrazinylidene]-2,3-dihydro, 1,1-dimethylethyl ester (CA INDEX NAME)

RN 855849-60-4 CAPLUS

CN Benzoic acid, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-61-5 CAPLUS

CN 2-Furancarboxylic acid, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-62-6 CAPLUS

CN 3-Pyridinecarboxylic acid, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-63-7 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-[4-(trifluoromethoxy)phenyl]- (CA INDEX NAME)

RN 855849-64-8 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-(2,4-dichlorophenyl)-(CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-65-9 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-(4-methoxyphenyl)-

(CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-66-0 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-(3-methoxyphenyl)- (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-67-1 CAPLUS

CN Hydrazinecarboxamide, N-(2-chlorophenyl)-2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]- (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-68-2 CAPLUS

CN Hydrazinecarboxamide, N-(3-chlorophenyl)-2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]- (CA INDEX NAME)

Double bond geometry as described by ${\tt E}$ or ${\tt Z}$.

RN 855849-69-3 CAPLUS

CN Hydrazinecarboxamide, N-(4-chlorophenyl)-2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]- (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-70-6 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)

RN 855849-71-7 CAPLUS

CN Benzoic acid, 4-(trifluoromethyl)-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-72-8 CAPLUS

CN Benzoic acid, 4-chloro-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-73-9 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-phenyl- (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-74-0 CAPLUS

CN Benzoic acid, 2-chloro-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-75-1 CAPLUS

CN Benzoic acid, 2-hydroxy-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-76-2 CAPLUS

CN Benzoic acid, 4-nitro-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-77-3 CAPLUS

CN Benzoic acid, 3,5-bis(trifluoromethyl)-,
2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-78-4 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-6-methoxy-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-79-5 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-6-methyl-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-80-8 CAPLUS

CN 3-Pyridinecarboxylic acid, 2-amino-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-81-9 CAPLUS

CN Hydrazinecarboxamide, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]-N-[4-(trifluoromethyl)phenyl]- (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-85-3 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-86-4 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[6-chloro-1'-[(2E)-3-(4-chlorophenyl)-2-propen-1-yl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

Double bond geometry as described by E or Z.

RN 855849-87-5 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[1'-[(4-chlorophenyl)methyl]-2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

RN 855849-94-4 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-[2,3-dihydro-1'-[[4-(trifluoromethyl)phenyl]methyl]spiro[1H-indene-1,4'-piperidin]-3-ylidene]hydrazide (CA INDEX NAME)

IT 855849-93-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of insecticidal spiroindane derivs. as insecticides, acaracides, molluscicides and nematocides)

RN 855849-93-3 CAPLUS

CN 4-Pyridinecarboxylic acid, 2-chloro-, 2-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-3-ylidene)hydrazide (CA INDEX NAME)

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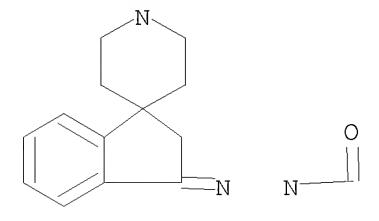
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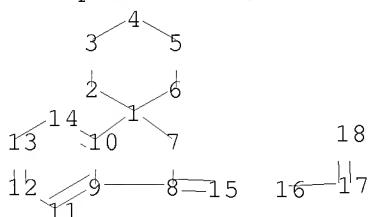
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CAS Information Use Policies apply and are available at:

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ring nodes :

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chain bonds :

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ring bonds :

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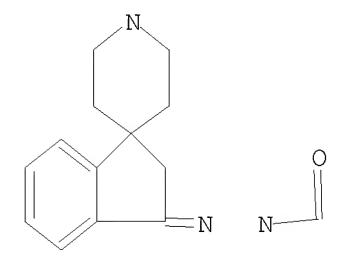
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L5 HAS NO ANSWERS

L5 STR



Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED 1291 ITERATIONS

5 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
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100.0% PROCESSED 26312 ITERATIONS

42 ANSWERS

SEARCH TIME: 00.00.01

L7 42 SEA SSS FUL L5

С

CAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2011.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

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L8 3 L7

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L8 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2011 ACS on STN GI

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 R^{7}
 R^{8}

Disclosed is an agent represented by a general formula I (A = CONHR1, ER2; AΒ R1 = (un) substituted C1-6 alkyl, (un) substituted C3-6 cycloalkyl; E =1,3,4-oxadiazol, 1,2,4-oxadiazol, imidazol; R2 = H, (un)substituted C1-6alkyl, phenyloxy, phenylamino, pyridylamino; R3, R4 = H, halogen, C1-6 alkyl, R5, R6, R7, R8 = H, halogen, C1-6 (halogenated) alkyl, C1-6 (halogenated)alkoxy; Z, Y = N, :CH- except Z = Y = :CH-; U = single bond, methylene, CH(OH), carbonyl, O, S, sulfinyl, sulfonyl; V = single bond, methylene, CH(OH); W = methylene, CH(OH), etc.; m, n = 0, 1 except m = 0/n= 1 and m = 1/n = 0), or is pharmaceutically acceptable salt as an active component. The agent has stearoyl-CoA desaturase-inhibitory effect, and suitable for use for prevention and/or treatment of related disease, e.g. obesity, hyperlipidemia, lipid metabolism disorder, diabetes, etc. For example, 6-(3,4-dihydro-1'H-spiro[chromene-2,4'-piperidin]-1'-yl)-N-(2hydroxy-2-phenylethyl)piridazine-3-carboxamide (II) was prepared The compound II showed ≥ 50 % stearoyl-CoA desaturase activity-inhibitory effect at 0.2 μ M in vitro. Also, a capsule containing II 50 mg/150 mg capsule was formulated.

AN 2009:1430440 CAPLUS

DN 151:565181

TI Pharmaceutical agent containing novel spiro piperidine derivatives having stearoyl-CoA desaturase-inhibitory effect

IN Uto, Yoshikazu; Kiyotsuka, Yohei

PA Daiichi Sankyo Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 136 pp. CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

P	ATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
	P 2009269850 P 2008-120937	A	20091119 20080507	JP 2008-120937	20080507	

OS MARPAT 151:565181

IT 1024604-69-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(novel spiro piperidine derivs. having stearoyl-CoA

desaturase-inhibitory effect)

RN 1024604-69-0 CAPLUS

CN 3-Pyridazinecarboxamide, 6-[2,3-dihydro-3-(methoxyimino)spiro[1H-indene-1,4'-piperidin]-1'-yl]-N-(2-hydroxy-2-phenylethyl)- (CA INDEX NAME)

OSC.G 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

The title compds. [I; A = CONHR1 or ER2; R1 = (un)substituted C1-6 alkylAB or C3-6 cycloalkyl; E = 1,3,4-oxadiazole, 1,2,4-oxadiazole, or imidazole group; R2 = H, each (un) substituted C1-6 alkyl, phenyloxy, phenylamino, or pyridylamino; R3, R4 = H, halo, C1-6 alkyl; R5-R8 = H, halo, C1-6 alkyl, C1-6 haloalkyl, C1-6 alkoxy, C1-6 haloalkoxy; Z, Y = N, CH; U = single bond, methylene, CH(OH), CH(OQ1), CO, C(:NOQ2), O; Q1, Q2 = C1-6 alkyl; m, n=0 or 1, provided that a case where m=0 and n=1 or m=1 and n=0is excluded] or pharmacol. acceptable salts were prepared These compds. have excellent stearoyl-CoA-desaturase inhibitory activity and are useful for the prevention and/or treatment of obesity, hyperlipidemia, hypertriglyceridemia, insulin resistance syndrome, abnormal glucose tolerance, diabetes, diabetes complications (including diabetes peripheral nerve disorder, diabetic nephropathy, diabetic retinopathy, and diabetic macroangiopathy), cataract, gestational diabetes, polycystic ovary syndrome, arteriosclerosis, atherosclerosis, diabetic arteriosclerosis, hypertension, cerebral vascular disorders, coronary artery disease, fatty liver, nonalcoholic fatty hepatitis, dyspnea, backache (lumbago), gonarthrosis, gout, and cholelithiasis. Thus, a suspension of 92 mg 6-chloro-N-(2-hydroxy-2-phenylethyl)pyridazine-3-carboxamide, 67 mg 3,4-Dihydro-1'H-spiro[chromene-2,4'-piperidine], and diisopropylethylamine in n-butanol was heated at 120° for 25 h to give 49% 6-(3,4-dihydrospiro[chromene-2,4'-piperidin]-1'-yl)-N-(2-hydroxy-2phenylethyl)pyridazine-3-carboxamide (II). II at 2 μ M inhibited ≥50% stearoyl-CoA-desaturase. A capsule and a tablet formulation containing II were described.

ΙI

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2008:583341 CAPLUS
AN
     148:538285
DN
     Preparation of spiropiperidine derivatives as stearoyl-CoA-desaturase
ΤI
     inhibitors
     Uto, Yoshikazu; Kiyotsuka, Yohei
IN
     Daiichi Sankyo Company, Limited, Japan
PA
     PCT Int. Appl., 233pp.
SO
     CODEN: PIXXD2
     Patent
DT
     Japanese
LA
FAN.CNT 1
     PATENT NO.
                         KIND
                                DATE
                                          APPLICATION NO.
                                                                    DATE
     WO 2008056687
                                20080515 WO 2007-JP71605
PΙ
                          Α1
                                                                    20071107
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA,
             CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI,
             GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG,
             KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME,
             MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL,
             PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
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             GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM
PRAI JP 2006-303866
                          Α
                                20061109
OS
     MARPAT 148:538285
     1024604-69-0P, N-(2-Hydroxy-2-phenylethyl)-6-[3-(methoxyimino)-
ΙT
     2,3-dihydrospiro[indene-1,4'-piperidin]-1'-yl]pyridazine-3-carboxamide
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (preparation of spiropiperidine derivs. as stearoyl-CoA-desaturase
        inhibitors)
     1024604-69-0 CAPLUS
RN
     3-Pyridazinecarboxamide, 6-[2,3-dihydro-3-(methoxyimino)spiro[1H-indene-
CN
     1,4'-piperidin]-1'-y1]-N-(2-hydroxy-2-phenylethy1)- (CA INDEX NAME)
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OSC.G 7 THERE ARE 7 CAPLUS RECORDS THAT CITE THIS RECORD (7 CITINGS)
RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2011 ACS on STN GI

$$R^{8}$$
 R^{9}
 R^{9

Title compds. I [X = O, amino; Y = bond, CO, CS, SOO-2; R1 = H, alkyl, alkoxycarbonyl, etc.; R2-3 = H, halo, CN, alkyl, etc.; R4 = halo, NO2, CN, etc.; Ra = H, halo, OH, CN, etc.; p, q = 0-6; R8 = alk(en/yn)yl, etc.] are prepared For instance, II is prepared in 3 steps from spiro[indan-1-one-3,4'-piperidine]-1'-carboxylic acid tert Bu ester, 4-chlorocinnamyl chloride and hydroxylamine (E (dominant) and Z oximes isolated). Selected example compds. gave >80% control of Spodoptera littoralis. I are useful in controlling insects, acarines, nematodes or molluscs.

AN 2005:570877 CAPLUS

DN 143:77964

TI Preparation of insecticidal spiroindane derivatives

IN Cassayre, Jerome; Molleyres, Louis-Pierre; Maienfisch, Peter; Cederbaum, Fredrik

PA Syngenta Participations A.-G., Switz.

SO PCT Int. Appl., 114 pp. CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

	PATENT NO.						D	DATE			APPL	DATE								
ΡI	WO	2005058836				 A1	20050630			WO 2004-IB4108						20041209				
	W: AE, AG, AL,			AM,	AT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,				
			CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,		
			GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KΖ,	LC,		
			LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,		
			NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,		
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			AZ,	BY,	KG,	KΖ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,		
			EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IS,	ΙΤ,	LT,	LU,	MC,	NL,	PL,	PT,		
			RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,		
			MR,	NE,	SN,	TD,	TG													
	EP 1697327			A1	20060906				EP 2004-806338						20041209					
	EP	1697	327			B1		2011	0713											
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,		

		I.	E, SI,	LT,	FI,	RO,	CY,	TR,	BG,	CZ,	EE,	, HU,	PL,	SK,	IS		
	BR	200401	7555		Α		2007	0327	B	R 2	2004-	-1755	5		,	2004120	19
	JP	200751	6253		${ m T}$		2007	0621	J]	2	2006-	-5436	59			2004120	19
	ΑT	516273			${ m T}$		2011	0715	A'	Γ 2	2004-	-8063	38			2004120	19
	ΙN	2006CN	02077		A		2007	0706	II	N 2	2006-	-CN20	77			2006061	. 2
	US	200803	06101		A1		2008	1211	U	S 2	2008-	-5811	77			2008082	28
PRAI	GB	2003-2	8906		Α		2003	1212									
	MO	2004-I	B4108		M		2004	1209									
ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT																	
OS	CAS	SREACT	143:77	964;	MARI	PAT	143:	77964	4								
IT	855	5849-47	-7P														
	RL:	: AGR (.	Agricu	ltura	al us	se);	BSU	(Bio	ologi	cal	l stı	udy,	uncl	lassit	fie	d); RCT	• •
	(Re	eactant); SPN	I (Syı	nthet	cic	prep	arat:	ion);	ВΙ	IOL	(Biol	ogic	cal st	cud	y); PRE	ŀΡ
	(Pr	reparat	ion);	RACT	(Rea	acta	int o	r rea	agent);	USES	S (Us	es)				
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		acarac						_									
RN	855	849-47	-7 CA	PLUS													
CN	Spi	lro[1H-	indene	-1, 4	'-pi	peri	dine] -1 '-	-carb	ЭХУ	ylic	acid	,				
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		l-dimet									_		_	•	_	·	
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RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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